MY MEDIC SPF 30 REEF FRIENDLY SUNSCREEN- homosalate, titanium dioxide, ethylhexyl salicylate, phenylbenzimidazole sulfonic acid cream Sled Distribution, LLC

81417-009 MY MEDIC SUNSCREEN SPF30

Active ingredient

Homosalate 14%

Titanium Dioxide 10%

Ethylhexyl Salicylate 4.5%

Phenylbenzimidazole Sulfonic Acid 1.5%

Purpose

Sunscreen

Uses

Helps prevent sunburn.

If used as directed with other sunburn protection measures (see Directions), decreases the risk of skin cancer and early skin aging caused by the sun.

Warnings

- For external use only
- **Do not use** on damaged or broken skin
- When using this product keep out of eyes.

Rinse with water to remove.

- Stop use and ask a doctor if rash occurs
- **Keep out of reach of children.** If product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Apply to skin liberally, avoiding the eye area.
- Wait at least 30 mins before sun exposure, or as directed by a physician.
- Re-apply after 80 mniutes of swimming and/or sweating.
- Re-apply immediately after towel drying and at least every 3 hours.

Inactive ingredients

Aluminum Hydroxide, Bisabolol, Butyloctyl Salicylate, Caprylhydroxamic Acid, Dimethicone, Glycerin, Glyceryl Caprylate, Glyceryl Stearate, Isohexadecane, Peg-100 Stearate, Polyacrylate Crosspolymer-6, Polyhydroxystearic Acid, Potassium Cetyl Phosphate, Propylene Glycol, Stearic Acid, Triethanolamine, VP Eicosene Copolymer, Water, Xanthan Gum, 1,2 Hexanediol



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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81417-009
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ENSULIZOLE (UNII: 9YQ9DI1W42) (ENSULIZOLE - UNII:9YQ9DI1W42)	ENSULIZOLE	1.5 g in 100 g	
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	4.5 g in 100 g	
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	14 g in 100 g	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	10 g in 100 g	

Inactive Ingredients	
Ingredient Name	Strength
GLYCERYL CAPRYLATE (UNII: TM2TZ D4G4A)	
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIMETHICONE (UNII: 92RU3N3Y10)	
PEG-100 STEARATE (UNII: YD01N1999R)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
XANTHAN GUM (UNII: TTV12P4NEE)	
LEVOMENOL (UNII: 24WE03BX2T)	
POTASSIUM CETYL PHOSPHATE (UNII: 03KCY6P7UT)	
CAPRYLHYDROXAMIC ACID (UNII: UPY805K99W)	
WATER (UNII: 059QF0KO0R)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE, DIMETHYLACRYLAMIDE, LAURYL METHACRYLATE AND LAURETH-4 METHACRYLATE COPOLYMER, TRIMETHYLOLPROPANE TRIACRYLATE CROSSLINKED (45000 MPA.S) (UNII: Q7UI015FF9)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
ISOHEXADECANE (UNII: 918X1OUF1E)	
POLYHYDROXYSTEARIC ACID STEARATE (UNII: 8KQ7I65XZE)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
VINYLPYRROLIDONE/EICOSENE COPOLYMER (UNII: 035MV9S1C3)	
TROLAMINE (UNII: 903K93S3TK)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:81417-009- 01	3.5 g in 1 POUCH; Type 0: Not a Combination Product	10/31/2022	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M020	10/31/2022		

Labeler - Sled Distribution, LLC (079772888)

Revised: 1/2024 Sled Distribution, LLC